| NITRATE-N + NITRITE-N IN DRINKING AND SURFACE WATERS, AND DOMESTIC AND INDUSTRIAL WASTES SEAL AQ2 METHOD NO: EPA 114 A REVISION 6 | | | | | | |
|--|--|---|-------------------|-----|----------|--|
| Facility Name: | VELAP ID | | | | | |
| Assessor Name: Analyst Name: | Analyst Name: | | Inspection Date | | | |
| Relevant Aspect of Standards | Method Reference | Y | N | N/A | Comments | |
| Records Examined: SOP Number/ Revision/ Date Analyst: | | | | | | |
| Sample ID: Date of Sample Prepa | D: Date of Sample Preparation: | | Date of Analysis: | | | |
| Is the linear calibration range determined initially, and does it contain a minimum of a blank and three standards? | Method Supplement 1, Rev. 2 (MS) 3.2.1 | | | | | |
| 2. Is linearity reestablished if any verification data exceeds initial calibration values by ±10%? | MS 3.2.1 | | | | | |
| 3. Is a laboratory control sample analyzed with every batch, and is recovery assessed against current laboratory criteria? NOTE: The laboratory "should" establish upper and lower control limits from control charts based on % recovery. | MS 3.4.3, 3.4.3.4, 3.4.3.5 | | | | | |
| 4. Is at least one method blank carried through all the procedural steps with each batch? | MS 3.4.1.1 | | | | | |
| 5. Is the calibration verified using a calibration standard after every ten samples or every analytical batch? | MS 4.5 | | | | | |
| Is a minimum of 10% of all samples spiked with the stock standard? | MS 3.3.1 | | | | | |
| 7. For compliance monitoring, is the concentration of the matrix spike at the regulatory limit OR 1 to 5 times higher than the background concentration of the sample? | MS 3.3.1.1.1 | | | | | |
| 8. Were absorbencies read at 520 nm? | 2.1 | | | | | |
| 9. Was volumetric glassware class A? | 6.2 | | | | | |
| 10. Was the pH of the ammonium chloride buffer stock adjusted to 8.5? | 7.1 | | | | | |
| 11. Did the working buffer contain 0.025% surfactant? | 7.1 | | | | | |
| 12. Was the working buffer discarded if it developed a pink color? | 7.1 | | | | | |
| 13. Was the sodium nitrate used to make stock Nitrate Standard solutions dried for at least 2 hours at 105°C? | 7.2 | | | | | |
| Notes/Comments: | | | | | | |

NITRATE-N + NITRITE-N IN DRINKING AND SURFACE WATERS, AND DOMESTIC AND INDUSTRIAL WASTES SEAL AQ2 METHOD NO: EPA-114-A REVISION 6 Υ N/A **Relevant Aspect of Standards** Method Ν Comments Reference Records Examined: SOP Number/ Revision/ Date _ Analyst:__ Sample ID: _ Date of Sample Preparation:___ _ Date of Analysis:__ 14. Were stock Nitrite Standard solutions stored in amber 7.2 bottles at 4°C for not longer than 1 month? 15. Were intermediate Nitrate Standard solutions stored 7.2 at 4°C for not longer than 2 weeks? 16. Were intermediate Nitrite Standard solutions prepared 7.2 at least twice weekly? 17. Were samples collected in glass or plastic bottles? 8.1 18. For Nitrate in unchlorinated Drinking Water samples: Were samples cooled to 4°C and analyzed within 48 8.2.1 hours? 19. For nitrate in drinking water, are samples preserved at 4°C and analyzed within 48 hours of collection 40CFR141.23.k(2) unless the sample is chlorinated? If chlorinated, analyze within 14 days. 20. For nitrite in drinking water, are samples preserved at 40CFR141.23.k(2) 4°C and analyzed within 48 hours? 21. For nitrate + nitrite in drinking water, are samples preserved by acidifying to pH<2 with sulfuric acid and 40CFR141.23.k(2) analyzed within 28 days? 22. For nitrate in nonpotable water, are samples 40CFR136.3 Table preserved at ≤6°C and analyzed within 48 hours? 23. For nitrite in nonpotable water, are samples 40CFR136.3 Table preserved at ≤6°C and analyzed within 48 hours? 24. For nitrate + nitrite in nonpotable water, are samples 40CFR136.3 Table acidified to pH<2 with sulfuric acid, preserved at ≤6°C, and analyzed within 28 days? Notes/Comments: